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November 29, 2004

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APPLICATION NUMBER: 60/511,353

FILING DATE: October 14, 2003

RELATED PCT APPLICATION NUMBER: PCT/US04/34446

Certified by



Jon W Dudas

Acting Under Secretary of Commerce
for Intellectual Property
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PTO/SB/17 (10-03)

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FEE TRANSMITTAL for FY 2004

Effective 10/01/2003. Patent fees are subject to annual revision.

☐ Applicant claims small entity status. See 37 CFR 1.27

TOTAL AMOUNT OF PAYMENT (\$) 160.00

Complete if Known

Application Number	UNKNOWN
Filing Date	HEREWITH
First Named Inventor	T.G. AMOS ET AL.
Examiner Name	UNKNOWN
Art Unit	UNKNOWN
Attorney Docket No.	CL2244 US PRV

15757
60/511353
101403**METHOD OF PAYMENT (check all that apply)**☐ Check ☐ Credit card ☐ Money Order ☐ Other ☐ None☒ Deposit Account:Deposit Account Number: 04-1928
Deposit Account Name: E. I. du Pont de Nemours and Company

The Director is authorized to: (check all that apply)

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☒ Charge any additional fee(s) or any underpayment of fee(s)
☐ Charge fee(s) indicated below, except for the filing fee to the above-identified deposit account.**FEE CALCULATION****1. BASIC FILING FEE**

Large Entity		Small Entity		Fee Description	Fee Paid
Fee Code	Fee (\$)	Fee Code	Fee (\$)		
1001	770	2001	385	Utility filing fee	
1002	340	2002	170	Design filing fee	
1003	530	2003	265	Plant filing fee	
1004	770	2004	385	Reissue filing fee	
1005	160	2005	80	Provisional filing fee	160.00
SUBTOTAL (1)				(\$)	160.00

2. EXTRA CLAIM FEES FOR UTILITY AND REISSUE

	Extra Claims	Fee from below	Fee Paid
Total Claims	-20** =	18	
Independent Claims	-3** =	86	
Multiple Dependent	YES	290.00	

Large Entity		Small Entity		Fee Description
Fee Code	Fee (\$)	Fee Code	Fee (\$)	
1202	18	2202	9	Claims in excess of 20
1201	86	2201	43	Independent claims in excess of 3
1203	290	2203	145	Multiple dependent claim, if not paid
1204	86	2204	43	** Reissue independent claims over original patent
1205	18	2205	9	** Reissue claims in excess of 20 and over original patent

SUBTOTAL (2) (\$) 0.00

**or number previously paid, if greater. For Reissues, see above

FEE CALCULATION (continued)**3. ADDITIONAL FEES**

Large Entity Small Entity

Fee Code	Fee (\$)	Fee Code	Fee (\$)	Fee Description	Fee Paid
1051	130	2051	65	Surcharge - late filing fee or oath	
1052	50	2052	25	Surcharge - late provisional filing fee or cover sheet	
1053	130	1053	130	Non-English specification	
1812	2,520	1812	2,520	For filing a request for ex parte reexamination	
1804	920*	1804	920*	Requesting publication of SIR prior to Examiner action	
1805	1,840*	1805	1,840*	Requesting publication of SIR after Examiner action	
1251	110	2251	55	Extension for reply within first month	
1252	420	2252	210	Extension for reply within second month	
1253	950	2253	475	Extension for reply within third month	
1254	1,480	2254	740	Extension for reply within fourth month	
1255	2,010	2255	1,005	Extension for reply within fifth month	
1401	330	2401	165	Notice of Appeal	
1402	330	2402	165	Filing a brief in support of an appeal	
1403	290	2403	145	Request for oral hearing	
1451	1,510	1451	1,510	Petition to institute a public use proceeding	
1452	110	2452	55	Petition to revive - unavoidable	
1453	1,330	2453	665	Petition to revive - unintentional	
1501	1,330	2501	665	Utility issue fee (or reissue)	
1502	480	2502	240	Design issue fee	
1503	640	2503	320	Plant issue fee	
1460	130	1460	130	Petitions to the Commissioner	
1807	50	1807	50	Processing fee under 37 CFR 1.17(q)	
1806	180	1806	180	Submission of Information Disclosure Stmt	
8021	40	8021	40	Recording each patent assignment per property (times number of properties)	
1809	770	2809	385	Filing a submission after final rejection (37 CFR 1.129(a))	
1810	770	2810	385	For each additional invention to be examined (37 CFR 1.129(b))	
1801	770	2801	385	Request for Continued Examination (RCE)	
1802	900	1802	900	Request for expedited examination of a design application	

Other fee (specify)

*Reduced by Basic Filing Fee Paid

SUBTOTAL (3) (\$) 0.00

(Completes if applicable)

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Date

OCTOBER 14, 2003

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PROVISIONAL APPLICATION FOR PATENT COVER SHEET

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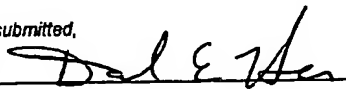
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Additional inventors are being named on the _____ separately numbered sheets attached hereto					
TITLE OF THE INVENTION (500 characters max)					
CHROMIUM OXIDE COMPOSITIONS CONTAINING ZINC, THEIR PREPARATION, AND THEIR USE AS CATALYSTS AND CATALYST PRECURSORS					
Direct all correspondence to: CORRESPONDENCE ADDRESS					
<input checked="" type="checkbox"/> Customer Number: 23906					
OR					
<input type="checkbox"/> Firm or Individual Name _____					
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Address _____					
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ENCLOSED APPLICATION PARTS (check all that apply)					
<input checked="" type="checkbox"/> Specification Number of Pages <u>26</u>					
<input checked="" type="checkbox"/> Drawing(s) Number of Sheets <u>2</u>					
<input type="checkbox"/> Application Date Sheet. See 37 CFR 1.76					
<input type="checkbox"/> CD(s), Number _____					
<input type="checkbox"/> Other (specify) 					
METHOD OF PAYMENT OF FILING FEES FOR THIS PROVISIONAL APPLICATION FOR PATENT					
<input type="checkbox"/> Applicant claims small entity status. See 37 CFR 1.27.					
<input type="checkbox"/> A check or money order is enclosed to cover the filing fees.					
<input checked="" type="checkbox"/> The Director is hereby authorized to charge filing fees or credit any overpayment to Deposit Account Number: <u>04-1928</u>					
<input type="checkbox"/> Payment by credit card. Form PTO-2038 is attached.					
FILING FEE Amount (\$) <div style="border: 1px solid black; padding: 10px; width: 100px; margin: 0 auto;">\$160.00</div>					
The invention was made by an agency of the United States Government or under a contract with an agency of the United States Government.					
<input checked="" type="checkbox"/> No.					
<input type="checkbox"/> Yes, the name of the U.S. Government agency and the Government contract number are: _____					

[Page 1 of 2]

Respectfully submitted,

SIGNATURE


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Date OCTOBER 14, 2003REGISTRATION NO. 31,366

(if appropriate)

Docket Number: CL2244 US PRV**USE ONLY FOR FILING A PROVISIONAL APPLICATION FOR PATENT**

This collection of information is required by 37 CFR 1.51. The information is required to obtain or retain a benefit by the public which is to file (and by the USPTO to process) an application. Confidentiality is governed by 35 U.S.C. 122 and 37 CFR 1.14. This collection is estimated to take 8 hours to complete, including gathering, preparing, and submitting the completed application form to the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, U.S. Department of Commerce, P.O. Box 1450, Alexandria, VA 22313-1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Mail Stop Provisional Application, Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450.

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PROVISIONAL APPLICATION COVER SHEET
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Docket Number CL2244 US PRV

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[Page 2 of 2]

Number _____ of _____

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TITLECHROMIUM OXIDE COMPOSITIONS CONTAINING ZINC, THEIR
PREPARATION, AND THEIR USE AS CATALYSTS AND CATALYST
PRECURSORS5 FIELD OF THE INVENTION

This invention relates to chromium-containing compositions, their preparation, and their use for the catalytic processing of hydrocarbons and/or halogenated hydrocarbons.

BACKGROUND

10 U. S. Patent No. 5,281,568 discloses a fluorination catalyst containing chromium and zinc. The amount of zinc may be in the range of from about 0.5% by weight to about 25% by weight.

Australian Patent Document No. AU-A-80340/94 discloses bulk or supported catalysts based on chromium oxide (or oxides of chromium) and
15 at least one other catalytically active metal (e.g., Mg, V, Mn, Fe, Co, Ni, or Zn), in which the major part of the oxide(s) is in the crystalline state (and when the catalyst is a bulk catalyst, its specific surface, after activation with HF, is at least 8 m²/g). The crystalline phases disclosed include Cr₂O₃, CrO₂, NiCrO₃, NiCrO₄, NiCr₂O₄, MgCrO₄, ZnCr₂O₄ and mixtures
20 of these oxides.

U. S. Patent No. 3,878,257 discloses the catalyzed reaction of 1,1,2-trichlorotrifluoropropene with hydrogen fluoride to form 2-chloropentafluoropropene in the presence of a catalyst combination of activated anhydrous chromium(III) oxide and a divalent zinc compound.

25 There remains a need for catalysts that can be used for processes such as the selective fluorination and chlorofluorination of saturated and unsaturated hydrocarbons, hydrochlorocarbons, hydrochlorofluorocarbons, and chlorofluorocarbons, the fluorination of unsaturated fluorocarbons, the isomerization and disproportionation of fluorinated organic compounds, the
30 dehydrofluorination of hydrofluorocarbons, and the chlorodefluorination of fluorocarbons.

SUMMARY OF THE INVENTION

This invention provides a chromium-containing catalyst composition comprising ZnCr₂O₄ (zinc chromite) and crystalline α -chromium oxide
35 wherein the ZnCr₂O₄ contains between about 10 atom percent and 67 atom percent of the chromium in the composition and at least about 70 atom percent of the zinc in the composition, and wherein at least about

90 atom percent of the chromium present as chromium oxide in the composition is present as ZnCr_2O_4 or crystalline α -chromium oxide.

This invention also provides a method for preparing said composition comprising ZnCr_2O_4 and crystalline α -chromium oxide. The method comprises (a) co-precipitating a solid by adding ammonium hydroxide (aqueous ammonia) to an aqueous solution of a soluble zinc salt and a soluble trivalent chromium salt that contains at least three moles of nitrate (i.e., NO_3^-) per mole of chromium (i.e., Cr^{3+}) in the solution and has a zinc concentration of from about 5 mole % to about 25 mole % of the total concentration of zinc and chromium in the solution and where at least three moles of ammonium (i.e., NH_4^+) per mole of chromium (i.e., Cr^{3+}) in the solution has been added to the solution, (b) collecting the co-precipitated solid formed in (a); (c) drying the collected solid; and (d) calcining the dried solid.

This invention also provides a chromium-containing catalyst composition, said composition being prepared by treatment of said composition comprising ZnCr_2O_4 and crystalline α -chromium oxide with a fluorinating agent (e.g., anhydrous hydrogen fluoride).

This invention also provides a process for changing the fluorine distribution (i.e., content and/or arrangement) in a halogenated hydrocarbon, or incorporating fluorine in a saturated or unsaturated hydrocarbon, in the presence of a catalyst. The process is characterized by using as a catalyst at least one composition selected from the group consisting of (i) the ZnCr_2O_4 and crystalline α -chromium oxide compositions of this invention and (ii) the ZnCr_2O_4 and crystalline α -chromium oxide compositions of this invention which have been treated with a fluorinating agent.

BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1 represents an energy dispersive spectroscopy spectrum of the zinc chromite phase present in a zinc/chromium oxide composition nominally containing 20 atom % zinc.

Figure 2 represents an energy dispersive spectroscopy spectrum of the α - Cr_2O_3 phase present in the same zinc/chromium oxide composition nominally containing 20 atom % zinc

DETAILED DESCRIPTION

The compositions of this invention are zinc- and chromium-containing oxides comprising ZnCr_2O_4 and crystalline α -chromium oxide wherein the ZnCr_2O_4 contains between about 10 atom percent and 67

atom percent of the chromium in the composition and at least about 70 atom percent of the zinc in the composition. The compositions disclosed include a catalytic composition comprising ZnCr_2O_4 and crystalline α -chromium oxide.

5 The compositions of this invention may be prepared by the method described above using co-precipitation followed by calcination. Such methods include, but are not limited to, precipitation from aqueous solutions containing chromium and zinc salts in the desired molar ratio.

10 In a typical co-precipitation technique, an aqueous solution of zinc and chromium(III) salts is prepared. The relative concentrations of the zinc and chromium(III) salts in the aqueous solution is dictated by the bulk atom percent zinc relative to chromium desired in the final catalyst. The concentration of zinc is from about 5 mole % to about 25 mole % of the total concentration of zinc and chromium in the solution. The
15 concentration of chromium(III) in the aqueous solution is typically in the range of 0.3 to 3 moles per liter with 0.75-1.5 moles per liter being a preferred concentration. While different chromium(III) salts might be employed, chromium(III) nitrate or its hydrated forms such as $[\text{Cr}(\text{NO}_3)_3(\text{H}_2\text{O})_9]$, are the most preferred chromium(III) salts for
20 preparation of said aqueous solution.

 While different zinc salts might be employed for preparation of said aqueous solutions, preferred zinc salts for preparation of catalysts for the process of this invention include zinc(II) nitrate and its hydrated forms such as $[\text{Zn}(\text{NO}_3)_2(\text{H}_2\text{O})_6]$.

25 The aqueous solution of the chromium(III) and zinc salts may then be evaporated either under vacuum or at elevated temperature to give a solid which is then calcined.

 It is preferred to treat the aqueous solution of the chromium(III) and zinc salts with a base such as ammonium hydroxide (aqueous ammonia)
30 to precipitate the zinc and chromium as the hydroxides. Bases containing alkali metals such as sodium or potassium hydroxide or the carbonates may be used but are not preferred. The addition of ammonium hydroxide to the aqueous solution of the chromium(III) and zinc salts is typically carried out gradually over a period of 1 to 12 hours. The pH of the solution
35 is monitored during the addition of base. The final pH is typically in the range of 6.0 to 11.0, preferably from about 7.5 to about 9.0, most preferably about 8.0 to 8.7. The precipitation of the zinc and chromium hydroxide mixture is typically carried out at a temperature of about 15°C to

about 60°C, preferably from about 20°C to about 40°C. After the ammonium hydroxide is added, the mixture is typically stirred for up to 24 hours. The precipitated chromium and zinc hydroxides serve as precursors to ZnCr_2O_4 and crystalline α -chromium oxide.

- 5 After the precipitation of the zinc and chromium hydroxide mixture is complete, the mixture is dried. This may be carried out by evaporation in an open pan on a hot plate or steam bath or in an oven or furnace at a suitable temperature. Suitable temperatures include temperatures from about 60°C to about 130°C (e.g., from about 100°C to about 120°C).
- 10 Alternatively the drying step may be carried out under vacuum using, for example, a rotary evaporator.

- Optionally, the precipitated zinc and chromium hydroxide mixture may be collected and, if desired, washed with deionized water before drying. Preferably the precipitated zinc and chromium hydroxide mixture is
- 15 not washed prior to the drying step.

- After the zinc and chromium hydroxide mixture has been dried, the nitrate salts are then decomposed by heating the solid from about 250°C to about 350°C. The resulting solid is then calcined at temperatures of from about 400°C to about 1000°C, preferably from about 400°C to about
- 20 900°C. The calcination may be carried out in a crucible or pan in an oven or furnace or in a tubular reactor. The calcination temperature can influence the activity of the catalysts and the product distribution. Lower calcination temperatures (i.e., those below about 500°C) may result in the presence of some residual nitrate impurities. The calcination is preferably
- 25 carried out in the presence of oxygen, most preferably in the presence of air.

 Of note are compositions comprising ZnCr_2O_4 and crystalline α -chromium oxide wherein the ZnCr_2O_4 is formed during the calcination step.

- 30 Of note are chromium-containing catalyst compositions of this invention which comprise ZnCr_2O_4 (zinc chromite) and crystalline α -chromium oxide wherein the ZnCr_2O_4 contains between about 20 atom percent and about 50 atom percent of the chromium in the composition. Also of note are chromium-containing catalyst compositions of this
- 35 invention which comprise ZnCr_2O_4 (zinc chromite) and crystalline α -chromium oxide wherein the ZnCr_2O_4 contains at least about 90 atom percent of the zinc in the composition. Also of note are chromium-containing catalyst compositions of this invention comprising zinc chromite

and crystalline α -chromium oxide wherein greater than 95 atom percent of the chromium that is not present as zinc chromite is present as crystalline α -chromium oxide. Also of note are chromium-containing catalyst compositions of this invention which consist essentially of ZnCr_2O_4 (zinc chromite) and crystalline α -chromium oxide.

The compositions of this invention may be characterized by well-established analytical techniques including transmission electron microscopy (TEM), energy dispersive spectroscopy (EDS) and X-ray diffraction. EDS is an analytical tool available in conjunction with scanning or analytical TEM.

The presence of zinc in the various zinc and chromium oxide compositions of this invention is clearly indicated by elemental analysis using EDS. EDS analyses of various Cr/Zn oxide samples having a Zn content of 2, 5, 10, and 20 atom % calcined at 900°C indicate the presence of two phases: a zinc chromite phase where the relative content of Cr to Zn is 2 to 1 and a chromium oxide phase with no evidence of zinc in the lattice (Zn is not detected in the EDS spectrum of this phase). For example, Figure 1 shows the EDS spectrum of the zinc chromite phase present in a zinc/chromium composition nominally containing 20 atom % zinc. For comparison, Figure 2 shows the EDS spectrum of the α - Cr_2O_3 phase present in the same zinc/chromium composition nominally containing 20 atom % zinc. In each of these Figures, X-ray intensity, I , representing thousands of counts is plotted against energy level, E , representing thousands of electron volts (keV). Peaks in each plot correlate with the presence of certain elements. Calibration experiments have shown that the relative heights of the $K\alpha$ peaks for the metals in mixed zinc and chromium oxide compositions reflect the mole ratio of zinc and chromium in the compositions. Thus, the EDS spectra are valid on a quantitative basis for elements whose atomic masses are fairly similar. These results are also consistent with wide-angle X-ray diffraction analysis experiments of the various samples, which indicated that each sample consisted of two phases: a pure α - Cr_2O_3 and a ZnCr_2O_4 spinel phase. There are no significant changes in the cell volume of the Cr_2O_3 phase in the Cr/Zn samples compared to zinc-free chromium oxide. This indicates that there is no detectable substitution of Zn in the Cr_2O_3 lattice. The weight percent of the ZnCr_2O_4 phase increases with increasing zinc concentration.

The compositions of this invention may further comprise one or more additives in the form of metal compounds that can alter the selectivity and/or activity of the catalyst compositions containing crystalline α -Cr₂O₃ and ZnCr₂O₄ or fluorinated α -Cr₂O₃ and ZnCr₂O₄. Suitable additives may be selected from the group consisting of fluorides, oxides, or oxyfluoride compounds of Mg, Ca, Zn, Sc, Y, La, Ti, Zr, Hf, V, Nb, Ta, Mo, W, Mn, Re, Fe, Ru, Co, Rh, Ir, Ni, Pd, Pt, Cu, Ag, Au, and Ce,

The total content of the additive(s) in the compositions of the present invention may be from about 0.05 atom % to about 15 atom % based on the total metal content of the compositions provided that when the additive is a compound of zinc, the additive comprises less than 30 mole percent of the total zinc in the final composition. The additives may be incorporated into the compositions of the present invention by standard procedures such as impregnation of a solution of the additive followed by drying or co-precipitation.

The calcined zinc chromite/ α -chromium oxide compositions of the present invention may be pressed into various shapes such as pellets for use in packing reactors. It may also be used in powder form.

Typically, the calcined compositions will be pre-treated with a fluorinating agent prior to use as catalysts for changing the fluorine content of halogenated carbon compounds. Typically this fluorinating agent is HF though other materials may be used such as sulfur tetrafluoride, carbonyl fluoride, and fluorinated carbon compounds such as trichlorofluoromethane, dichlorodifluoromethane, chlorodifluoromethane, trifluoromethane, or 1,1,2-trichlorotrifluoroethane. This pretreatment can be accomplished, for example, by placing the catalyst in a suitable container which can be the reactor to be used to perform the process of the instant invention, and thereafter, passing HF over the dried, calcined catalyst so as to partially saturate the catalyst with HF. This is conveniently carried out by passing HF over the catalyst for a period of time, for example, about 0.1 to about 10 hours at a temperature of, for example, about 200°C to about 450°C. Nevertheless, this pretreatment is not essential.

As noted above catalysts provided in accordance with this invention may be used for changing the fluorine distribution and/or content of halogenated hydrocarbons. The catalysts of this invention may also be used to incorporate fluorine into a saturated or unsaturated hydrocarbon. Processes for changing the fluorine distribution in halogenated

hydrocarbons include fluorination, chlorofluorination, isomerization, disproportionation, dehydrofluorination and chlorodefluorination.

Processes for incorporating fluorine into saturated or unsaturated hydrocarbons include chlorofluorination of saturated or unsaturated

- 5 hydrocarbons and fluorination of unsaturated hydrocarbons. The processes of this invention are characterized by using as a catalyst at least one composition selected from the group consisting of the ZnCr_2O_4 / α -chromium oxide compositions of this invention and the ZnCr_2O_4 / α -chromium oxide compositions of this invention which have been treated
- 10 with a fluorinating agent. Of note are processes wherein the fluorine content of a halogenated hydrocarbon compound or an unsaturated hydrocarbon compound is increased by reacting said compound with hydrogen fluoride in the vapor phase in the presence of said catalyst composition. Also of note are processes wherein the fluorine content of a
- 15 halogenated hydrocarbon compound or a hydrocarbon compound is increased by reacting said compound with HF and Cl_2 in the vapor phase in the presence of said catalyst composition. Also of note are processes wherein the fluorine distribution in a halogenated hydrocarbon compound is changed by isomerizing said halogenated hydrocarbon compound in the
- 20 presence of said catalyst composition. Also of note are processes wherein the fluorine distribution in a halogenated hydrocarbon compound is changed by disproportionating said halogenated hydrocarbon compound in the vapor phase in the presence of said catalyst composition. Also of note are processes wherein the fluorine content of a halogenated
- 25 hydrocarbon compound is decreased by dehydrofluorinating said halogenated hydrocarbon compound in the presence of said catalyst composition. Also of note are processes wherein the fluorine content of a halogenated hydrocarbon compound is decreased by reacting said halogenated hydrocarbon compound with hydrogen chloride in the vapor
- 30 phase in the presence of said catalyst composition.

- Typical of saturated halogenated hydrocarbons suitable for fluorination, chlorofluorination, isomerization, disproportionation, dehydrofluorination and chlorodefluorination processes are those which have the formula $\text{C}_n\text{H}_a\text{Br}_b\text{Cl}_c\text{F}_d$, wherein n is an integer from 1 to 6, a is
- 35 an integer from 0 to 12, b is an integer from 0 to 4, c is an integer from 0 to 13, d is an integer from 0 to 13, the sum of b, c and d is at least 1 and the sum of a, b, c, and d is equal to $2n + 2$, provided that n is at least 2 for isomerization, disproportionation and dehydrofluorination processes, a is

at least one for dehydrofluorination processes, b is 0 for chlorodefluorination processes, b + c is at least 1 for fluorination processes and is 0 for dehydrofluorination processes, a + b + c is at least 1 for fluorination, chlorofluorination, isomerization, disproportionation and dehydrofluorination processes and d is at least 1 for isomerization, disproportionation, dehydrofluorination and chlorodefluorination processes. Typical of saturated hydrocarbon compounds suitable for chlorofluorination are those which have the formula C_qH_r where q is an integer from 1 to 6 and r is $2q + 2$. Typical of unsaturated halogenated hydrocarbons suitable for fluorination, chlorofluorination, isomerization, disproportionation, and chlorodefluorination processes are those which have the formula $C_pH_eBr_fCl_gF_h$, wherein p is an integer from 2 to 6, e is an integer from 0 to 10, f is an integer from 0 to 2, g is an integer from 0 to 12, h is an integer from 0 to 11, the sum of f, g and h is at least 1 and the sum of e, f, g, and h is equal to $2p$, provided that f is 0 for chlorodefluorination processes, e + f + g is at least 1 for isomerization and disproportionation processes and h is at least 1 for isomerization, disproportionation and chlorodefluorination processes. Typical of saturated hydrocarbons suitable for chlorofluorination are those which have the formula C_qH_r where q is an integer from 1 to 6 and r is $2q + 2$. Typical of unsaturated hydrocarbons suitable for fluorination and chlorofluorination are those which have the formula C_iH_j where i is an integer from 2 to 6 and j is $2i$.

The fluorination and chlorofluorination processes are typically conducted in the vapor phase in a tubular reactor at temperatures of from about 150°C to 500°C. For saturated compounds the fluorination is preferably carried out from about 175°C to 400°C and more preferably from about 200°C to about 350°C. For unsaturated compounds the fluorination is preferably carried out from about 150°C to 350°C and more preferably from about 175°C to about 300°C. The reactions are typically conducted at atmospheric and superatmospheric pressures. For reasons of convenience in downstream separations processes (e.g., distillation), pressures of up to about 30 atmospheres may be employed. The contact time in the reactor is typically from about 1 to about 120 seconds and preferably from about 5 to about 60 seconds.

The amount of HF reacted with the unsaturated hydrocarbons or halogenated hydrocarbon compounds should be at least a stoichiometric amount. The stoichiometric amount is based on the number of Br and/or Cl substituents to be replaced by F in addition to one mole of HF to

saturate the carbon-carbon double bond if present. Typically, the molar ratio of HF to the said compounds of the formulas $C_nH_aBr_bCl_cF_d$, $C_pH_eBr_fCl_gF_h$, and C_iH_j can range from about 0.5:1 to about 100:1, preferably from about 2:1 to about 50:1, and more preferably from about 3:1 to about 20:1. In general, with a given catalyst composition, the higher the temperature and the longer the contact time, the greater is the conversion to fluorinated products. The above variables can be balanced, one against the other, so that the formation of higher fluorine substituted products is maximized.

Examples of saturated compounds of the formula $C_nH_aBr_bCl_cF_d$ which may be reacted with HF in the presence of the catalyst of this invention include CH_2Cl_2 , CH_2Br_2 , $CHCl_3$, CCl_4 , C_2Cl_6 , C_2BrCl_5 , C_2Cl_5F , $C_2Cl_4F_2$, $C_2Cl_3F_3$, $C_2Cl_2F_4$, C_2ClF_5 , C_2HCl_5 , C_2HCl_4F , $C_2HCl_3F_2$, $C_2HCl_2F_3$, C_2HClF_4 , C_2HBrF_4 , $C_2H_2Cl_4$, $C_2H_2Cl_3F$, $C_2H_2Cl_2F_2$, $C_2H_2ClF_3$, $C_2H_3Cl_3$, $C_2H_3Cl_2F$, $C_2H_3ClF_2$, $C_2H_4Cl_2$, C_2H_4ClF , $C_3Cl_6F_2$, $C_3Cl_5F_3$, $C_3Cl_4F_4$, $C_3Cl_3F_5$, C_3HCl_7 , C_3HCl_6F , $C_3HCl_5F_2$, $C_3HCl_4F_3$, $C_3HCl_3F_4$, $C_3HCl_2F_5$, $C_3H_2Cl_6$, $C_3H_2BrCl_5$, $C_3H_2Cl_5F$, $C_3H_2Cl_4F_2$, $C_3H_2Cl_3F_3$, $C_3H_2Cl_2F_4$, $C_3H_2ClF_5$, $C_3H_3Cl_5$, $C_3H_3Cl_4F$, $C_3H_3Cl_3F_2$, $C_3H_3Cl_2F_3$, $C_3H_3ClF_4$, $C_3H_4Cl_4$, $C_4Cl_4Cl_4$, $C_4Cl_4Cl_6$, $C_4H_5Cl_5$, $C_4H_5Cl_4F$, and $C_5H_4Cl_8$.

Specific examples of fluorination reactions of saturated halogenated hydrocarbon compounds which may be carried out under the conditions described above using the catalysts of this invention include the conversion of CH_2Cl_2 to CH_2F_2 , the conversion of $CHCl_3$ to a mixture of $CHCl_2F$, $CHClF_2$, and CHF_3 , the conversion of CH_3CHCl_2 to a mixture of CH_3CHClF and CH_3CHF_2 , the conversion of CH_2ClCH_2Cl to a mixture of CH_3CHClF and CH_3CHF_2 , the conversion of CH_3CCl_3 to a mixture of CH_3CCl_2F , CH_3CClF_2 , and CH_3CF_3 , the conversion of CH_2ClCF_3 to CH_2FCF_3 , the conversion of $CHCl_2CF_3$ to a mixture of $CHClFCF_3$ and CHF_2CF_3 , the conversion of $CHClFCF_3$ to CHF_2CF_3 , the conversion of $CHBrFCF_3$ to CHF_2CF_3 , the conversion of $CCl_3CF_2CCl_3$ to a mixture of $CCl_2FCF_2CClF_2$ and $CClF_2CF_2CClF_2$, the conversion of $CCl_3CH_2CCl_3$ to $CF_3CH_2CF_3$ or a mixture of $CF_3CH_2CClF_2$ and $CF_3CH_2CF_3$, the conversion of $CCl_3CH_2CHCl_2$ to a mixture of $CF_3CH_2CHF_2$, $CF_3CH=CHCl$, and $CF_3CH=CHF$, the conversion of $CF_3CCl_2CClF_2$ to a mixture of $CF_3CCl_2CF_3$, and $CF_3CClFCF_3$, the conversion of $CF_3CCl_2CF_3$ to $CF_3CClFCF_3$, and the conversion of a mixture comprising

CF₃CF₂CHCl₂ and CClF₂CF₂CHClF to a mixture of CF₃CF₂CHClF and CF₃CF₂CHF₂.

Examples of unsaturated compounds of the formula C_pH_eBr_fCl_gF_h and C_iH_j which may be reacted with HF in the presence of the catalysts of this invention include C₂Cl₄, C₂BrCl₃, C₂Cl₃F, C₂Cl₂F₂, C₂ClF₃, C₂F₄, C₂HCl₃, C₂HBrCl₂, C₂HCl₂F, C₂HClF₂, C₂HF₃, C₂H₂Cl₂, C₂H₂ClF, C₂H₂F₂, C₂H₃Cl, C₂H₃F, C₂H₄, C₃H₆, C₃H₅Cl, C₃H₄Cl₂, C₃H₃Cl₃, C₃H₂Cl₄, C₃HCl₅, C₃Cl₆, C₃Cl₅F, C₃Cl₄F₂, C₃Cl₃F₃, C₃Cl₂F₄, C₃ClF₅, C₃HF₅, C₃H₂F₄, C₃F₆, C₄Cl₈, C₄Cl₂F₆, C₄ClF₇, C₄H₂F₆, and C₄HClF₆.

Specific examples of fluorination reactions of unsaturated halogenated hydrocarbon compounds which may be carried out using the catalysts of this invention include the conversion of CHCl=CCl₂ to a mixture of CH₂ClCF₃ and CH₂FCF₃, the conversion of CCl₂=CCl₂ to a mixture of CHCl₂CF₃, CHClFCF₃, and CHF₂CF₃, the conversion of CCl₂=CH₂ to a mixture of CH₃CCl₂F, CH₃CClF₂, and CH₃CF₃, the conversion of CH₂=CHCl to a mixture of CH₃CHClF and CH₃CHF₂, the conversion of CF₂=CH₂ to CH₃CF₃, the conversion of CCl₂=CClCF₃ to a mixture of CF₃CCl=CF₂ and CF₃CHClCF₃, the conversion of CF₃CF=CF₂ to CF₃CHFCF₃, the conversion of CF₃CH=CF₂ to CF₃CH₂CF₃, and the conversion of CF₃CH=CHF to CF₃CH₂CHF₂.

Also of note is a catalytic process for producing a mixture of 2-chloro-1,1,3,3,3-pentafluoropropene (i.e., CF₃CCl=CF₂ or CFC-1215xc) and 2-chloro-1,1,1,3,3,3-hexafluoropropane (i.e., CF₃CHClCF₃ or HCFC-226da) by the fluorination of one or more halopropene compounds CX₃CCl=CClX, wherein each X is independently selected from the group consisting of F and Cl. Preferred halopropenes of the formula CX₃CCl=CClX include 1,2,2-trichloro-3,3,3-trifluoro-1-propene (i.e., CCl₂=CClCF₃ or CFC-1213xa) and hexachloropropene (i.e., CCl₂=CClCCl₃). The CFC-1215xc/HCFC-226da mixture is produced by reacting the above unsaturated compound(s) with HF in the vapor phase in the presence of the catalysts of this invention at temperatures from about 240°C to about 400°C, preferably about 250°C to about 350°C.

The amount of HF fed to the reactor should be at least a stoichiometric amount based on the number of Cl substituents in the CX₃CCl=CClX starting material(s). In the case of fluorination of CFC-1213xa, the stoichiometric ratio of HF to CFC-1213xa is 3:1 for synthesis of HCFC-226da. Preferred ratios of HF to CX₃CCl=CClX starting material(s) are typically in the range of about twice the stoichiometric ratio

to about 30:1. Preferred contact times are from 1 to 60 seconds. In contrast to catalyst compositions comprising chromium oxide in the absence of zinc, the catalyst compositions of this invention provide mixtures of CFC-1215xc and HCFC-226da.

5 Further information on the fluorination of CFC-1213xa to a mixture of CFC-1215xc and HCFC-226da is provided in U.S. Patent Application _____ [CL2372 US PRV] filed concurrently herewith, and hereby incorporated by reference herein in its entirety.

10 Mixtures of saturated halogenated hydrocarbon compounds or mixtures of unsaturated hydrocarbons and/or halogenated hydrocarbon compounds may also be used in the vapor phase fluorination reactions as well as mixtures comprising both unsaturated hydrocarbons and halogenated hydrocarbon compounds. Specific examples of mixtures of saturated halogenated hydrocarbon compounds and mixtures of
15 unsaturated hydrocarbons and unsaturated halogenated hydrocarbon compounds that may be subjected to vapor phase fluorination using the catalysts of this invention include a mixture of CH_2Cl_2 and $\text{CCl}_2=\text{CCl}_2$, a mixture of $\text{CCl}_2\text{FCClF}_2$ and CCl_3CF_3 , a mixture of $\text{CCl}_2=\text{CCl}_2$ and $\text{CCl}_2=\text{CClCCl}_3$, a mixture of $\text{CH}_2=\text{CHCH}_3$ and $\text{CH}_2=\text{CClCH}_3$, a mixture of
20 CH_2Cl_2 and CH_3CCl_3 , a mixture of $\text{CHF}_2\text{CClF}_2$ and CHClFCF_3 , a mixture of $\text{CHCl}_2\text{CCl}_2\text{CH}_2\text{Cl}$ and $\text{CCl}_3\text{CHClCH}_2\text{Cl}$, a mixture of $\text{CHCl}_2\text{CH}_2\text{CCl}_3$ and $\text{CCl}_3\text{CHClCH}_2\text{Cl}$, a mixture of $\text{CHCl}_2\text{CHClCCl}_3$, $\text{CCl}_3\text{CH}_2\text{CCl}_3$, and $\text{CCl}_3\text{CCl}_2\text{CH}_2\text{Cl}$, a mixture of $\text{CHCl}_2\text{CH}_2\text{CCl}_3$ and $\text{CCl}_3\text{CH}_2\text{CCl}_3$, a mixture of $\text{CF}_3\text{CH}_2\text{CCl}_2\text{F}$ and $\text{CF}_3\text{CH}=\text{CCl}_2$, and a mixture of
25 $\text{CF}_3\text{CH}=\text{CHCl}$ and $\text{CF}_3\text{CH}=\text{CCl}_2$.

Where chlorine (Cl_2) is present as in chlorofluorinations, the amount of chlorine fed to the reactor is based on whether the halogenated hydrocarbon compounds fed to the reactor is unsaturated and the number of hydrogens in $\text{C}_n\text{H}_a\text{Br}_b\text{Cl}_c\text{F}_d$, C_qH_r , $\text{C}_p\text{H}_e\text{Br}_f\text{Cl}_g\text{F}_h$, and C_iH_j that are to
30 be replaced by chlorine and fluorine. One mole of Cl_2 is required to saturate a carbon-carbon double bond and a mole of Cl_2 is required for each hydrogen to be replaced by chlorine or fluorine. A slight excess of chlorine over the stoichiometric amount may be necessary for practical reasons, but large excesses of chlorine will result in complete
35 chlorofluorination of the products. The ratio of Cl_2 to halogenated carbon compound is typically from about 1:1 to about 10:1.

Specific examples of vapor phase chlorofluorination reactions of saturated halogenated hydrocarbon compounds of the general formula

$C_nH_aBr_bCl_cF_d$ and saturated hydrocarbon compounds of the general formula C_qH_r which may be carried out using the catalysts of this invention include the conversion of C_2H_6 to a mixture containing CH_2ClCF_3 , the conversion of CH_2ClCF_3 to a mixture of $CHClCF_3$ and CHF_2CF_3 , the
 5 conversion of $CCl_3CH_2CH_2Cl$ to a mixture of $CF_3CCl_2CClF_2$, $CF_3CCl_2CF_3$, $CF_3CClFCClF_2$, and $CF_3CClFCF_3$, the conversion of $CCl_3CH_2CHCl_2$ to a mixture of $CF_3CCl_2CClF_2$, $CF_3CCl_2CF_3$, $CF_3CClFCClF_2$, and $CF_3CClFCF_3$, the conversion of $CCl_3CHClCH_2Cl$ to a mixture of $CF_3CCl_2CClF_2$, $CF_3CCl_2CF_3$, $CF_3CClFCClF_2$, and
 10 $CF_3CClFCF_3$, the conversion of $CHCl_2CCl_2CH_2Cl$ to a mixture of $CF_3CCl_2CClF_2$, $CF_3CCl_2CF_3$, $CF_3CClFCClF_2$, and $CF_3CClFCF_3$, the conversion of $CCl_3CH_2CH_2Cl$ to a mixture of $CF_3CCl_2CHF_2$, $CF_3CClFCHF_2$, $CF_3CClFCClF_2$, and $CF_3CCl_2CF_3$, and the conversion of $CCl_3CH_2CHCl_2$ to a mixture of $CF_3CCl_2CHF_2$, $CF_3CClFCHF_2$,
 15 $CF_3CClFCClF_2$, and $CF_3CCl_2CF_3$.

Specific examples of vapor phase chlorofluorination reactions of unsaturated halogenated hydrocarbon compounds of the general formula $C_pH_eBr_fCl_gF_h$ and unsaturated hydrocarbon compounds of the general formula C_iH_j which may be carried out using the catalysts of this invention
 20 include the conversion of C_2H_4 to a mixture of CCl_3CClF_2 , CCl_2FCCl_2F , $CClF_2CCl_2F$, CCl_3CF_3 , CF_3CCl_2F , and $CClF_2CClF_2$, the conversion of C_2Cl_4 to a mixture of CCl_3CClF_2 , CCl_2FCCl_2F , $CClF_2CCl_2F$, CCl_3CF_3 , CF_3CCl_2F , and $CClF_2CClF_2$, and the conversion of C_3H_6 or $CF_3CCl=CCl_2$ to a mixture of $CF_3CCl_2CClF_2$, $CF_3CCl_2CF_3$, $CF_3CClFCClF_2$, and
 25 $CF_3CClFCF_3$.

Of note is a catalytic process for producing a mixture of 1,2,2-trichloro-1,1,3,3,3-pentafluoropropane (i.e., $CClF_2CCl_2CF_3$ or CFC-215aa) and 1,1,2-trichloro-1,2,3,3,3-pentafluoropropane (i.e., $CF_3CClFCCl_2F$ or CFC-215bb), by the chlorofluorination of a halopropene
 30 of the formula $CX_3CCl=CClX$, wherein each X is independently selected from the group F and Cl. Preferred halopropenes of the formula $CX_3CCl=CClX$ include 1,2,2-trichloro-3,3,3-trifluoro-1-propene (i.e., $CCl_2=CClCF_3$ or CFC-1213xa) and hexachloropropene (i.e., $CCl_2=CClCCl_3$). The mixture of CFC-215aa and CFC-215bb is produced
 35 by reacting the above unsaturated compounds with Cl_2 and HF in the vapor phase in the presence of the catalysts of this invention at temperatures from about 200°C to about 400°C, preferably about 250°C to 350°C.

Further information on the chlorofluorination of CFC-1213xa to produce CFC-215aa and CFC-215bb is provided in U.S. Patent Application _____ [CL2320 US PRV] filed concurrently herewith, and hereby incorporated by reference herein in its entirety.

5 Also of note is a catalytic process for producing a mixture of 2,2-dichloro-1,1,1,3,3,3-hexafluoropropane (i.e., $\text{CF}_3\text{CCl}_2\text{CF}_3$ or CFC-216aa) and 1,2-dichloro-1,1,2,3,3,3-hexafluoropropane (i.e., $\text{CF}_3\text{CClFCClF}_2$ or CFC-216ba), by the chlorofluorination of a halopropene of the formula $\text{CX}_3\text{CCl}=\text{CX}_2$, wherein each X is independently selected
10 from the group F and Cl. Preferred halopropenes of the formula $\text{CX}_3\text{CCl}=\text{CClX}$ include 1,2,2-trichloro-3,3,3-trifluoro-1-propene (i.e., $\text{CCl}_2=\text{CClCF}_3$ or CFC-1213xa) and hexachloropropene (i.e., $\text{CCl}_2=\text{CClCCl}_3$). The mixture of CFC-216aa and CFC-216ba is produced by reacting the above unsaturated compounds with Cl_2 and HF in the
15 vapor phase in the presence of the catalysts of this invention at temperatures from about 230°C to about 425°C, preferably about 250°C to 400°C.

Further information on the chlorofluorination of CFC-1213xa to produce CFC-216aa and CFC-216ba is provided in U.S. Patent
20 Application _____ [CL2246 US PRV] filed concurrently herewith, and hereby incorporated by reference herein in its entirety.

Mixtures of saturated hydrocarbon compounds and saturated halogenated hydrocarbon compounds and mixtures of unsaturated hydrocarbon compounds and unsaturated halogenated hydrocarbon
25 compounds as well as mixtures comprising both saturated and unsaturated compounds may be chlorofluorinated using the catalysts of the present invention. Specific examples of mixtures of saturated and unsaturated hydrocarbons and halogenated hydrocarbons that may be used include a mixture of $\text{CCl}_2=\text{CCl}_2$ and $\text{CCl}_2=\text{CClCCl}_3$, a mixture of
30 $\text{CHCl}_2\text{CCl}_2\text{CH}_2\text{Cl}$ and $\text{CCl}_3\text{CHClCH}_2\text{Cl}$, a mixture of $\text{CHCl}_2\text{CH}_2\text{CCl}_3$ and $\text{CCl}_3\text{CHClCH}_2\text{Cl}$, a mixture of $\text{CHCl}_2\text{CHClCCl}_3$, $\text{CCl}_3\text{CH}_2\text{CCl}_3$, and $\text{CCl}_3\text{CCl}_2\text{CH}_2\text{Cl}$, a mixture of $\text{CHF}_2\text{CH}_2\text{CF}_3$ and $\text{CHCl}=\text{CHCF}_3$, and a mixture of $\text{CH}_2=\text{CH}_2$ and $\text{CH}_2=\text{CHCH}_3$.

Included in the present invention are embodiments in which the
35 fluorine distribution of a halogenated hydrocarbon compound is changed by rearranging the H, Br, Cl, and F substituents in the molecule (typically to a thermodynamically preferred arrangement) while maintaining the

same number of the H, Br, Cl, and F substituents, respectively. This process is referred to herein as isomerization.

5 In another embodiment of the present invention, the fluorine distribution of a halogenated hydrocarbon compound is changed by exchanging at least one F substituent of one molecule of the halogenated hydrocarbon starting material with at least one H, Br and/or Cl substituent of another molecule of the halogenated hydrocarbon starting material so as to result in the formation of one or more halogenated hydrocarbon compounds having a decreased fluorine content compared to the
10 halogenated hydrocarbon starting material and one or more halogenated hydrocarbon compounds having an increased fluorine content compared to the halogenated hydrocarbon starting material. This process is referred to herein as disproportionation.

15 In another embodiment of the present invention, both isomerization and disproportionation reactions may occur simultaneously.

Whether carrying out isomerization, disproportionation or both isomerization and disproportionation, the fluorine distribution of saturated compounds of the formula $C_nH_aBr_bCl_cF_d$ and/or unsaturated compounds of the formula $C_pH_eBr_fCl_gF_h$ may be changed in the presence of a catalyst
20 composition of this invention.

The isomerization and disproportionation reactions are typically conducted at temperatures of from about 150°C to 500°C, preferably from about 200°C to about 400°C. The contact time in the reactor is typically from about 1 to about 120 seconds and preferably from about 5 to about
25 60 seconds. The isomerization and disproportionation reactions may be carried out in the presence of an inert gas such as helium, argon, or nitrogen. The isomerization and disproportionation reactions may be carried out in the presence of HF and HCl.

Specific examples of vapor phase isomerization reactions which
30 may be carried out using the catalysts of this invention include the conversion of $CClF_2CCl_2F$ to CCl_3CF_3 , the conversion of $CClF_2CClF_2$ to CF_3CCl_2F , the conversion of CHF_2CClF_2 to CF_3CHClF , the conversion of CHF_2CHF_2 to CF_3CH_2F , the conversion of $CF_3CClFCClF_2$ to $CF_3CCl_2CF_3$, and the conversion of $CF_3CHFCHF_2$ to $CF_3CH_2CF_3$.

35 Specific examples of vapor phase disproportionation reactions which may be carried out using the catalysts of this invention include the conversion of $CClF_2CClF_2$ to a mixture of $CClF_2CCl_2F$, CCl_3CF_3 , and

CF_3CClF_2 , and the conversion of CHClFCH_3 to a mixture of CHCl_2CF_3 , and CHF_2CF_3 .

Of note is a process for the conversion of a mixture of 2-chloro-1,1,2,2-tetrafluoroethane (i.e., $\text{CHF}_2\text{CClF}_2$ or HCFC-124a) and 2-chloro-1,1,1,2-tetrafluoroethane (i.e., CF_3CHClF or HCFC-124) to a mixture comprising 2,2-dichloro-1,1,1-trifluoroethane (i.e., CHCl_2CF_3 or HCFC-123) and 1,1,1,2,2-pentafluoroethane (i.e., CF_3CHF_2 or HFC-125) in addition to unconverted starting materials. The mixture comprising HFC-125 and HCFC-123 may be obtained in the vapor phase by contacting a mixture of HCFC-124a and HCFC-124 over the catalysts of this invention optionally in the presence of a diluent selected from the group consisting of HF, HCl, nitrogen, helium, argon, and carbon dioxide. The disproportionation is preferably conducted at about 150°C to about 400°C, more preferably about 250°C to about 350°C. If used, the diluent gas may be present in a molar ratio of diluent to haloethane of from about 1:1 to about 5:1. Preferred contact times are from about 10 seconds to about 60 seconds.

Included in this invention is a process for decreasing the fluorine content of a halogenated hydrocarbon compound by dehydrofluorinating said halogenated hydrocarbon compound in the presence of the catalyst of this invention. Halogenated hydrocarbon compounds suitable as starting materials for the dehydrofluorination processes of this invention include those of the general formula $\text{C}_n\text{H}_a\text{F}_d$, wherein n is an integer from 2 to 6, a is an integer from 1 to 12, d is an integer from 1 to 13, and the sum of a and d is equal to $2n + 2$.

The dehydrofluorination reactions are typically conducted at temperatures of from about 200°C to about 500°C, preferably from about 300°C to about 450°C. The contact time in the reactor is typically from about 1 to about 360 seconds and preferably from about 5 to about 120 seconds. The dehydrofluorination reactions can also be carried out in the presence of an inert gas such as helium, argon, or nitrogen to increase the extent of dehydrofluorination of the halogenated hydrocarbon compound.

The product of dehydrofluorination reaction consists of HF and the unsaturated fluorinated carbon compound resulting from loss of HF from the starting material. Specific examples of vapor phase dehydrofluorination reactions which may be carried out using the catalysts of this invention include the conversion of CH_3CHF_2 to $\text{CH}_2=\text{CHF}$, the

conversion of CH_3CF_3 to $\text{CH}_2=\text{CF}_2$, the conversion of $\text{CF}_3\text{CH}_2\text{F}$ to $\text{CF}_2=\text{CHF}$, the conversion of $\text{CHF}_2\text{CH}_2\text{CF}_3$ to $\text{CHF}=\text{CHCF}_3$, and the conversion of $\text{CF}_3\text{CH}_2\text{CF}_3$ to $\text{CF}_3\text{CH}=\text{CF}_2$.

Of note is a catalytic process for producing fluoroethene (i.e., $\text{CH}_2=\text{CHF}$ or vinyl fluoride) by the dehydrofluorination of a 1,1-difluoroethane (i.e., CHF_2CH_3 or HFC-152a). A mixture comprising vinyl fluoride and unconverted HFC-152a may be obtained in the vapor phase by contacting HFC-152a over the catalysts of this invention optionally in the presence of a diluent selected from the group consisting of HF, nitrogen, helium, argon, and carbon dioxide. The dehydrofluorination is preferably conducted at about 150°C to about 400°C , more preferably about 250°C to about 350°C . If used, the diluent gas may be present in a molar ratio of diluent to haloethane of from about 1:1 to about 5:1. Preferred contact times are from about 10 seconds to about 60 seconds.

Also of note is a catalytic process for producing 1,1,3,3,3-pentafluoropropene (i.e., $\text{CF}_2=\text{CHCF}_3$ or HFC-1225zc) by the dehydrofluorination of 1,1,1,3,3,3-hexafluoropropane (i.e., $\text{CF}_3\text{CH}_2\text{CF}_3$ or HFC-236fa). A mixture comprising HFC-1225zc and unconverted HFC-236fa may be obtained in the vapor phase by contacting HFC-236fa over the catalysts of this invention optionally in the presence of a diluent selected from the group consisting of HF, nitrogen, helium, argon, and carbon dioxide. The dehydrofluorination is preferably conducted at about 250°C to about 450°C , more preferably about 300°C to about 400°C . If used, the diluent gas may be present in a molar ratio of diluent to haloethane of from about 1:1 to about 5:1. Preferred contact times are from about 10 seconds to about 60 seconds.

Included in this invention is a process for decreasing the fluorine content of a halogenated hydrocarbon compound by reacting said halogenated hydrocarbon compound with hydrogen chloride (HCl) in the vapor phase in the presence of a catalyst composition of this invention. Such a process is referred to herein as a chlorodefluorination. Chlorodefluorination is disclosed in U.S. Patent No. 5,345,017 and U.S. Patent No. 5,763,698, which two patents are hereby incorporated herein by reference.

Halogenated hydrocarbon compounds suitable as starting materials for the chlorodefluorination processes of this invention may be saturated or unsaturated. Saturated halogenated hydrocarbon compounds suitable for the chlorodefluorination processes of this invention include those of the

general formula $C_nH_aCl_cF_d$, wherein n is an integer from 1 to 6, a is an integer from 0 to 12, c is an integer from 0 to 13, d is an integer from 1 to 13, and the sum of a, c and d is equal to $2n + 2$. Unsaturated halogenated hydrocarbon compounds suitable for the chlorodefluorination processes of this invention include those of the general formula $C_pH_eCl_gF_h$, wherein p is an integer from 2 to 6, e is an integer from 0 to 10, g is an integer from 0 to 12, h is an integer from 1 to 11, and the sum of e, g, and h is equal to $2p$. The fluorine content of saturated compounds of the formula $C_nH_aCl_cF_d$ and/or unsaturated compounds of the formula $C_pH_eCl_gF_h$ may be decreased by reacting said compounds with HCl in the vapor phase in the presence of the catalyst compositions of this invention.

The product of chlorodefluorination reactions typically comprise unreacted HCl, HF, unconverted starting material, saturated halogenated hydrocarbon compounds having a lower fluorine content than the starting material and unsaturated halogenated compounds. Specific examples of vapor phase chlorodefluorination reactions which may be carried out using the catalysts of this invention include the conversion of CHF_3 to a mixture of $CHCl_3$, $CHCl_2F$, and $CHClF_2$, the conversion of $CClF_2CClF_2$ to a mixture of CCl_3CCl_3 , CCl_3CCl_2F , CCl_3CClF_2 , CCl_2FCCl_2F , $CClF_2CCl_2F$, and CCl_3CF_3 , the conversion of CF_3CClF_2 to a mixture of CCl_3CCl_3 , CCl_3CCl_2F , CCl_3CClF_2 , CCl_2FCCl_2F , $CClF_2CCl_2F$, CCl_3CF_3 , $CClF_2CClF_2$, and CF_3CCl_2F , the conversion of $CF_3CCl_2CF_3$ to a mixture of $CF_3CCl_2CClF_2$, $CF_3CCl_2CCl_2F$, $CF_3CCl_2CCl_3$, and $CClF_2CCl_2CCl_3$, and the conversion of $CF_3CH_2CF_3$ to a mixture of $CCl_2=CHCF_3$, and $CCl_2=CClCF_3$.

The reaction products obtained by the processes of this invention can be separated by conventional techniques, such as with combinations including, but not limited to, scrubbing, decantation, or distillation. Some of the products of the various embodiments of this invention may form one or more azeotropes with each other or with HF.

The reactor, distillation columns, and their associated feed lines, effluent lines, and associated units used in applying the process of this invention should be constructed of materials resistant to hydrogen fluoride and hydrogen chloride. Typical materials of construction, well-known to the fluorination art, include stainless steels, in particular of the austenitic type, the well-known high nickel alloys, such as MonelTM nickel-copper alloys, HastelloyTM nickel-based alloys and, InconelTM nickel-chromium alloys, and copper-clad steel.

The processes of this invention can be carried out readily using well known chemical engineering practices.

Utility

Several of the reaction products obtained through use of the catalysts disclosed herein will have desired properties for direct commercial use. For example, CH_2F_2 (HFC-32), CHF_2CF_3 (HFC-125), CHF_2CF_3 (HFC-125), CH_2FCHF_2 (HFC-134), $\text{CF}_3\text{CH}_2\text{CF}_3$ (HFC-236fa), and $\text{CF}_3\text{CH}_2\text{CHF}_2$ (HFC-245fa) find application as refrigerants, CH_2FCF_3 (HFC-134a) and $\text{CF}_3\text{CHF}_2\text{CF}_3$ (HFC-227ea) find application as propellants, CH_2FCHF_2 (HFC-134) and $\text{CF}_3\text{CH}_2\text{CHF}_2$ (HFC-245fa) find application as blowing agents, and CHF_2CF_3 (HFC-125), $\text{CF}_3\text{CH}_2\text{CF}_3$ (HFC-236fa), and $\text{CF}_3\text{CHF}_2\text{CF}_3$ (HFC-227ea) find application as fire extinguishants.

Other reaction products obtained through the use of this invention are used as chemical intermediates to make useful products. For example, CCl_3CF_3 (CFC-113a) can be used to prepare CFC-114a which can then be converted to CH_2FCF_3 (HFC-134a) by hydrodechlorination. Similarly, $\text{CF}_3\text{CCl}_2\text{CF}_3$ (CFC-216aa) and $\text{CF}_3\text{CHClCF}_3$ (HCFC-226da) can be used to prepare $\text{CF}_3\text{CH}_2\text{CF}_3$ (HFC-236fa) by hydrodechlorination. Also, $\text{CF}_3\text{CCl}=\text{CF}_2$ (CFC-1215xc) and $\text{CF}_3\text{CCl}_2\text{CClF}_2$ (CFC-215aa) can be used to prepare $\text{CF}_3\text{CH}_2\text{CHF}_2$ (HFC-245fa) by hydrogenation and $\text{CF}_3\text{CClFCClF}_2$ (CFC-216ba) can be used to prepare $\text{CF}_3\text{CF}=\text{CF}_2$ (HFP).

The following specific embodiments are to be construed as merely illustrative, and do not constrain the remainder of the disclosure in any way whatsoever.

EXAMPLES

LEGEND

CFC-113 is $\text{CClF}_2\text{CCl}_2\text{F}$	CFC-114 is $\text{CClF}_2\text{CClF}_2$
CFC-114a is $\text{CF}_3\text{CCl}_2\text{F}$	HCFC-124 is CF_3CHClF
HCFC-124a is $\text{CClF}_2\text{CHF}_2$	HFC-125 is CF_3CHF_2
CFC-133a is $\text{CF}_3\text{CH}_2\text{Cl}$	226da is $\text{CF}_3\text{CHClCF}_3$
227ea is $\text{CF}_3\text{CHF}_2\text{CF}_3$	236fa is $\text{CF}_3\text{CH}_2\text{CF}_3$
HCC-1110 is $\text{CCl}_2=\text{CCl}_2$	CFC-1111 is $\text{CClF}=\text{CCl}_2$
HCC-1120 is $\text{CHCl}=\text{CCl}_2$	HCFC-1121 is $\text{CHCl}=\text{CClF}$
1215xc is $\text{CF}_3\text{CCl}=\text{CF}_2$	1225zc is $\text{CF}_3\text{CH}=\text{CF}_2$

Catalyst Characterization

Energy Dispersive Spectroscopy (EDS) and Transmission Electron Microscopy (TEM)

In these studies, the crystallites were analyzed using a Philips CM-20 high-resolution transmission electron microscope operated at an accelerating voltage of 200 kV and configured with an Oxford windowless EDS system with a Si(Li) elemental detector. In the EDS analyses, electron-transparent thin sections of samples were used to minimize sample thickness effects such as fluorescence. Also, due to the similarity of their atomic masses, the X-ray absorption cross-sections for Cr and Zn were assumed to be the same (see the discussion by Zaluzec on pages 121 to 167 in *Introduction to Analytical Electron Microscopy* edited by J. J. Hren, J. I. Goldstein, and D. C. Joy (Plenum Press, New York, 1979). The presence of copper in the EDS is due to the TEM grid and background in the microscope.

X-ray Powder Diffraction (XRD)

X-ray diffraction measurements were collected with a Philips XPERT automated powder diffractometer, Model 3040, using CuK(alpha) radiation ($\lambda = 1.5406 \text{ \AA}$). These measurements involved a scan between $2-90^\circ 2\text{-theta}$, with a step of 0.03° and a count time of 2 seconds. The volume dimensions and weight percents were determined using GSAS software and the Rietveld method. The volume average crystallite sizes were estimated using the Materials Data Jade software.

Catalyst Preparation

PREPARATION EXAMPLE 1

Preparation of 95%Chromium/5% Zinc Catalyst (450°C)

A solution of 380.14 g $\text{Cr}(\text{NO}_3)_3[9(\text{H}_2\text{O})]$ (0.950 mole) and 14.87 g $\text{Zn}(\text{NO}_3)_2[6(\text{H}_2\text{O})]$ (0.050 mole) was prepared in 1000 mL of deionized water. The solution was treated with 450 mL of 7.4M aqueous ammonium hydroxide over the course of one hour; the pH increased from 1.7 to pH 8.4. The slurry was stirred at room temperature overnight and then dried at 120°C in an oven in the presence of air. The dried solid was then calcined in air at 450°C for 20 hours; the resulting solid weighed 76.72 g.

PREPARATION EXAMPLE 2

Preparation of 90% Chromium/10% Zinc Catalyst (900°C)

A solution of 360.13 g $\text{Cr}(\text{NO}_3)_3[9(\text{H}_2\text{O})]$ (0.900 mole) and 29.75 g $\text{Zn}(\text{NO}_3)_2[6(\text{H}_2\text{O})]$ (0.100 mole) was prepared in 1000 mL of deionized water. The solution was treated with 450 mL of 7.4M aqueous ammonium

hydroxide over the course of 1.4 hours; the pH increased from 1.9 to pH 8.4. The slurry was stirred at room temperature overnight and then dried at 120°C in the presence of air. The dried solid was then calcined in air at 900°C for 20 hours; the resulting solid weighed 75.42 g.

5 X-ray powder analysis of the sample indicated the presence of three phases: the silicon internal standard, Cr_2O_3 (eskolaite), and ZnCr_2O_4 (zinc chromite). The weight % ZnCr_2O_4 was determined to be 23.9%. There was no significant change in the cell volume of the Cr/Zn sample (0.2896 nm^3) in comparison with a 900°C-calcined Cr_2O_3 sample
10 which had been precipitated in the absence of zinc (0.2895 nm^3). This indicates that zinc had not been substituted into the Cr_2O_3 lattice. The estimated coherent domain size of the Cr_2O_3 and ZnCr_2O_4 phases was 814 angstroms and 712 angstroms, respectively.

Analysis of the sample by TEM and EDS indicated the presence of
15 chromium oxide phases containing zinc having a Cr/Zn ratio of 2 and a chromium oxide phase containing no zinc.

PREPARATION EXAMPLE 3

Preparation of 95%Chromium/5% Zinc Catalyst (900°C)

A solution of 380.14 g $\text{Cr}(\text{NO}_3)_3[9(\text{H}_2\text{O})]$ (0.950 mole) and 14.87 g
20 $\text{Zn}(\text{NO}_3)_2[6(\text{H}_2\text{O})]$ (0.050 mole) was prepared in 1000 mL of deionized water. The solution was treated with 450 mL of 7.4M aqueous ammonium hydroxide over the course of one hour; the pH increased from 1.7 to pH 8.4. The slurry was stirred at room temperature overnight and then dried at 120°C in an oven in the presence of air. The dried solid was then
25 calcined in air at 900°C for 20 hours; the resulting solid weighed 70.06 g.

X-ray powder analysis of the sample indicated the presence of three phases: the silicon internal standard, Cr_2O_3 (eskolaite), and ZnCr_2O_4 (zinc chromite). The weight % ZnCr_2O_4 was determined to be 12.1%. There was no significant change in the cell volume of the Cr/Zn
30 sample (0.2894 nm^3) in comparison with a 900°C-calcined Cr_2O_3 sample which had been precipitated in the absence of zinc (0.2895 nm^3). This indicates that zinc had not been substituted into the Cr_2O_3 lattice. The estimated coherent domain size of the Cr_2O_3 and ZnCr_2O_4 phases was 962 angstroms and 913 angstroms, respectively.

35 Analysis of the sample by TEM and EDS indicated the presence of chromium oxide phases containing zinc having a Cr/Zn ratio of 2 and a chromium oxide phase containing no zinc.

PREPARATION EXAMPLE 4

Preparation of 80%Chromium/20% Zinc Catalyst (900°C)

A solution of 320.12 g of $\text{Cr}(\text{NO}_3)_3[9(\text{H}_2\text{O})]$ (0.800 mole) and 59.49 g $\text{Zn}(\text{NO}_3)_2[6(\text{H}_2\text{O})]$ (0.200 mole) was prepared in 1000 mL of deionized water. The solution was treated with 450 mL of 7.4M aqueous ammonium hydroxide over the course of one hour; the pH increased from about 1.7 to about pH 8.4. The slurry was stirred at room temperature overnight and then dried at 120°C in an oven in the presence of air. The dried solid was then calcined in air at 900°C for 22 hours; the resulting solid weighed 75.80 g.

X-ray powder analysis of the sample indicated the presence of three phases: the silicon internal standard, Cr_2O_3 (eskolaite), and ZnCr_2O_4 (zinc chromite). The weight % ZnCr_2O_4 was determined to be 60.9%. There was no significant change in the cell volume of the Cr/Zn sample (0.2896 nm^3) in comparison with a 900°C-calcined Cr_2O_3 sample which had been precipitated in the absence of zinc (0.2895 nm^3). This indicates that zinc had not been substituted into the Cr_2O_3 lattice. The estimated coherent domain size of the Cr_2O_3 and ZnCr_2O_4 phases was 779 angstroms and 679 angstroms, respectively.

Analysis of the sample by TEM and EDS indicated the presence of chromium oxide phases containing zinc having a Cr/Zn ratio of 2 and a chromium oxide phase containing no zinc.

PREPARATION EXAMPLE 5

Preparation of 98.1% Chromium/1.9% Zinc Catalyst (550°C)

A solution of 516.46 g $\text{Cr}(\text{NO}_3)_3[9(\text{H}_2\text{O})]$ (1.29 moles) and 7.31 g $\text{Zn}(\text{NO}_3)_2[6(\text{H}_2\text{O})]$ (0.0246 mole) was prepared in 500 mL of distilled water in 1L beaker resting on a hot plate. The mixture was then transferred to a Pyrex™ container and the container placed in a furnace. The container was heated from room temperature to 125°C at 10°C/min and then held at 125°C for six hours. The container was heated from 125°C to 350°C at 1°C/min and then held at 350°C for six hours. The container was heated from 350°C to 550°C at 1°C/min and then held at 550°C for 24 hours.

EXAMPLE 1

$\text{CF}_3\text{CH}_2\text{CF}_3$ Dehydrofluorination

A sample of the catalyst prepared in PREPARATION EXAMPLE 5, was pelletized (-12 to +20 mesh, (1.68 to 0.84 mm)); 29.86 g (20 mL) and sieved, and placed in a 5/8" (1.58 cm) diameter Inconel™ nickel alloy reactor tube heated in a fluidized sand bath. The catalyst was purged with

nitrogen at about 250°C prior to use. In addition, the catalyst, which had been previously used in chlorofluorination reactions, was originally fluorinated using a 1:1 ratio of HF and nitrogen (50 cc/min (8.3(10)⁻⁷ m³/sec) each) at 175°C. The catalyst was then treated with a 4:1 ratio of HF and nitrogen (nitrogen flow rate: 20 cc/min (3.3(10)⁻⁷ m³/sec); HF flow rate: 80 cc/min (1.3(10)⁻⁶ m³/sec) as the reactor temperature was gradually increased from 175°C to 400°C over several hours. HFC-236fa and nitrogen were fed to the reactor in a 1:4 molar ratio with a catalyst contact time of 15 seconds at a nominal pressure of one atmosphere. The GC-MS analyses of the reactor effluent at 300°C and 400°C are given below.

		Mole %	
		<u>300°C</u>	<u>400°C</u>
	<u>Component</u>		
	HFC-236fa	92.3	59.7
	HFC-1225zc	6.8	36.2
15	HFC-227ea	0.6	3.1
Minor products included CH ₃ CF ₃ , C ₃ F ₈ , C ₄ HF ₇ , CF ₃ CHClCF ₃ , and C ₃ HCIF ₄			

EXAMPLE 2

Dehydrofluorination of a CF₃CHClCF₃/CF₃CH₂CF₃ Mixture

A sample of the catalyst prepared in PREPARATION EXAMPLE 2 was pelletized (-12 to +20 mesh, (1.68 to 0.84 mm)); 26.64 g (15 mL) and sieved and placed in a 5/8" (1.58 cm) diameter Inconel™ nickel alloy reactor tube heated in a fluidized sand bath. The catalyst was purged with nitrogen at about 250°C prior to use. In addition, the catalyst, which had been previously used in chlorofluorination reactions, was originally fluorinated following a procedure similar to that in EXAMPLE 1. Nitrogen and a mixture comprising HCFC-226da (73.9%), HFC-236fa (25.5%), HFC-1225zc (0.2%), and CFC-216aa (0.1%) were co-fed to the reactor with catalyst contact time of 30 seconds at a nominal pressure of one atmosphere. The molar ratio of nitrogen to fluoropropane mixture was 4:1. The GC-MS analyses of the reactor effluent at 300°C and 400°C are given below.

		GC Area %	
		<u>300°C</u>	<u>400°C</u>
		<u>Component</u>	
	HCFC-226da	73.2	51.4
35	HFC-236fa	24.7	15.3
	HFC-1225zc	1.5	11.3
	CFC-1215xc	0.2	17.0

Minor products included CH_3CF_3 , $\text{C}_3\text{H}_3\text{F}_3$, $\text{C}_3\text{H}_2\text{F}_4$, $\text{CF}_3\text{CF}=\text{CHF}$, $\text{CF}_3\text{CHClCF}_3$, C_3HClF_4 , $\text{C}_3\text{Cl}_2\text{F}_6$, and $\text{C}_3\text{Cl}_2\text{F}_4$.

EXAMPLE 3

Disproportionation of a $\text{CF}_3\text{CHClF}/\text{CClF}_2\text{CHF}_2$ Mixture

- 5 Nitrogen and a mixture comprising HCFC-124a (97.1 mole %), HCFC-124 (2.4 mole %), and CFC-114 (0.4 mole %) were co-fed to the reactor containing the catalyst used in EXAMPLE 2. The molar ratio of nitrogen to the 124/124a mixture was 2:1 and the contact time was 30 seconds. The GC-MS analyses of the reactor effluent at 300°C and 400°C are given
10 below.

		Mole %	
		<u>300°C</u>	<u>400°C</u>
	<u>Component</u>		
	HFC-125	1.5	21.4
	HCFC-124	2.1	3.7
15	HCFC-124a	94.9	61.9
	CFC-133a	-	2.1
	$\text{C}_2\text{HCl}_2\text{F}_3$ isomers	0.7	3.3
	HCC-1110	-	0.5
	HCC-1120	-	1.4
20	CFC-1111	0.03	2.2
	$\text{C}_2\text{Cl}_2\text{F}_2$ isomers	0.06	1.7
	HCFC-1121	0.02	0.7
	CFC-114a	-	2.1
	CFC-114	0.4	0.4
25	CFC-113	-	0.3

Minor products included $\text{CF}_3\text{CH}_2\text{CF}_3$, $\text{CF}_3\text{CH}_2\text{Cl}$, C_3HF_5 , $\text{CF}_3\text{CHClCF}_3$, $\text{C}_2\text{HCl}_3\text{F}_2$, $\text{C}_2\text{Cl}_2\text{F}_4$, C_2ClF_5 and $\text{C}_2\text{Cl}_2\text{F}_2$.

CLAIMS

What is claimed is:

1. A chromium-containing catalyst composition, comprising:
ZnCr₂O₄; and
5 crystalline α -chromium oxide;
wherein the ZnCr₂O₄ contains between about 10 atom percent
and 67 atom percent of the chromium in the composition and at least
about 70 atom percent of the zinc in the composition, and wherein at least
about 90 atom percent of the chromium present as chromium oxide in the
10 composition is present as ZnCr₂O₄ or crystalline α -chromium oxide.
2. The chromium-containing catalyst composition of Claim 1
wherein the ZnCr₂O₄ contains between about 20 atom percent and about
50 atom percent of the chromium in the composition.
3. The chromium-containing catalyst composition of Claim 1
15 wherein the ZnCr₂O₄ contains at least about 90 atom percent of the zinc in
the composition.
4. The chromium-containing catalyst composition of Claim 1
wherein greater than 95% of the chromium that is not present as zinc
chromite is present as crystalline α -chromium oxide.
- 20 5. The chromium-containing catalyst composition of Claim 1 which
consists essentially of ZnCr₂O₄ and crystalline α -chromium oxide.
6. A chromium-containing catalyst composition prepared by
treatment of the composition of Claim 1 with a fluorinating agent.
7. The chromium-containing catalyst composition of Claim 6
25 wherein the fluorinating agent is anhydrous hydrogen fluoride.
8. A process for changing the fluorine distribution in a halogenated
hydrocarbon, or incorporating fluorine in a saturated or unsaturated
hydrocarbon, in the presence of a catalyst characterized by: using as a
catalyst at least one composition selected from the group consisting of
30 (i) the chromium-containing catalyst compositions of Claim 1 and
(ii) chromium-containing catalyst compositions prepared by treatment of a
composition of Claim 1 with a fluorinating agent.
9. The process of Claim 8 wherein the fluorine content of a
halogenated hydrocarbon compound or an unsaturated hydrocarbon
35 compound is increased by reacting said compound with hydrogen fluoride
in the vapor phase in the presence of said catalyst composition.
10. The process of Claim 8 wherein the fluorine content of a
halogenated hydrocarbon compound or a hydrocarbon compound is

increased by reacting said compound with HF and Cl₂ in the vapor phase in the presence of said catalyst composition.

11. The process of Claim 8 wherein the fluorine distribution in a halogenated hydrocarbon compound is changed by isomerizing said halogenated hydrocarbon compound in the presence of said catalyst composition.

12. The process of Claim 8 wherein the fluorine distribution in a halogenated hydrocarbon compound is changed by disproportionating said halogenated hydrocarbon compound in the vapor phase in the presence of said catalyst composition.

13. The process of Claim 8 wherein the fluorine content of a halogenated hydrocarbon compound is decreased by dehydrofluorinating said halogenated hydrocarbon compound in the presence of said catalyst composition.

14. The process of Claim 8 wherein the fluorine content of a halogenated hydrocarbon compound is decreased by reacting said halogenated hydrocarbon compound with hydrogen chloride in the vapor phase in the presence of said catalyst composition.

15. A method for preparing the chromium-containing catalyst composition of Claim 1, comprising:

- (a) co-precipitating a solid by adding ammonium hydroxide to an aqueous solution of a soluble zinc salt and a soluble trivalent chromium salt that contains at least three moles of nitrate per mole of chromium in the solution and has a zinc concentration of from about 5 mole % to about 25 mole % of the total concentration of zinc and chromium in the solution and where at least three moles of ammonium per mole of chromium in the solution has been added to the solution;
- (b) collecting the co-precipitated solid formed in (a);
- (c) drying the collected solid; and
- (d) calcining the dried solid.

16. The process of Claim 15 wherein ZnCr₂O₄ is formed during (d).

TITLECHROMIUM OXIDE COMPOSITIONS CONTAINING ZINC, THEIR
PREPARATION, AND THEIR USE AS CATALYSTS AND CATALYST
PRECURSORS

5

ABSTRACT OF THE DISCLOSURE

A chromium-containing catalyst is disclosed which includes both ZnCr_2O_4 and crystalline α -chromium oxide. The ZnCr_2O_4 contains between about 10 atom percent and 67 atom percent of the chromium in the composition and at least about 70 atom percent of the zinc in the composition, and at least about 90 atom percent of the chromium present as chromium oxide in the composition is present as ZnCr_2O_4 or crystalline α -chromium oxide. Also disclosed are a method for preparing this composition comprising ZnCr_2O_4 and crystalline α -chromium oxide; and a chromium-containing catalyst composition prepared by treatment of the composition comprising ZnCr_2O_4 and crystalline α -chromium oxide with a fluorinating agent. Also disclosed is a process for changing the fluorine distribution in a halogenated hydrocarbon, or incorporating fluorine in a saturated or unsaturated hydrocarbon, in the presence of at least one composition selected from the group consisting of (i) the ZnCr_2O_4 and crystalline α -chromium oxide compositions and (ii) the ZnCr_2O_4 and crystalline α -chromium oxide compositions which have been treated with a fluorinating agent.

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DEH/dmm

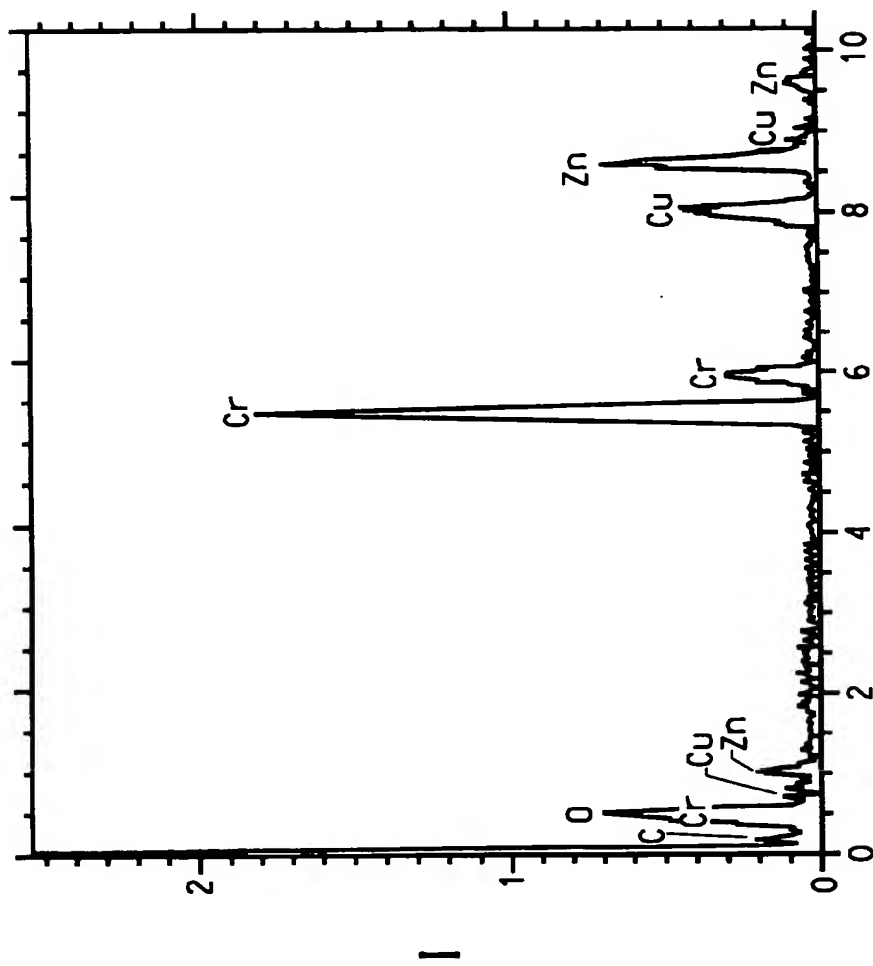


FIG. 1

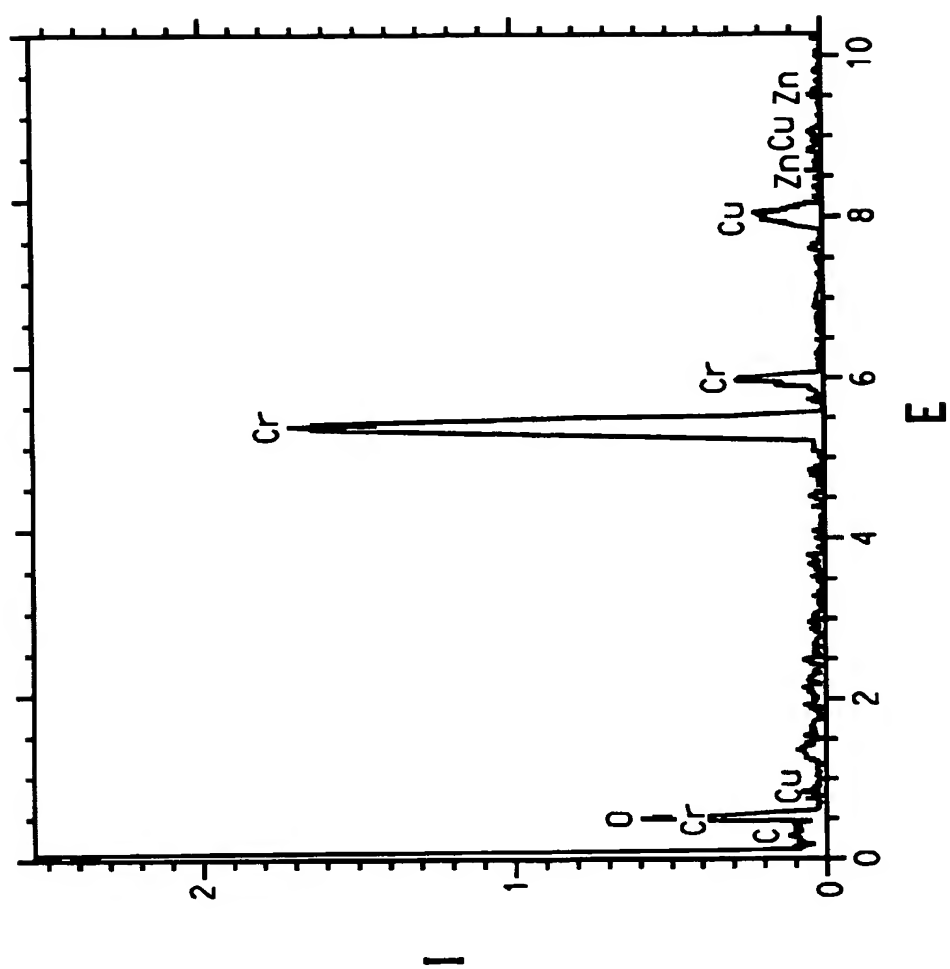


FIG. 2

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